2025-2026

CERTIFICATE OF ELECTRONIC DRUG LISTING

This certifies that:

AQUA MEDICA, S.A. DE C.V.
Carretera Federal México Cuautla Km. 65.8 No. 8
Tetelcingo, Cuautla 62757
México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**AQUACID-100 Acid Concentrate for Hemodialysis (3,785 mL)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-605-01**
U.S. Agent:	**Latin FDA Registrar**

Latin FDA Registrar will confirm that such listing remains effective upon request and presentation of this certificate until the end of the year shown above, unless terminated after issuance of this certificate. Latin FDA Registrar makes no other representations of warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Latin FDA Registrar assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Latin FDA Registrar waives, and will hold harmless and indemnify Latin FDA Registrar from any and all claims in connection with this product, its labeling, FDA drug listing, commerce or use. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of drug listing or the assignment of an DNC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue certificates of drug listing, nor does the U.S. Food and Drug Administration recognize certificates of drug listing. Latin FDA Registrar is not affiliated with the U.S. Food and Drug Administration.

Registro FDA

3942 West 11th Place, Inglewood CA. 90303
Telephone: +1-424-789-3544
info@fdaregistro.com

César López
Executive Director
Latin FDA Registrar Corp.
Dated: November 05, 2025

2025-2026

CERTIFICATE OF ELECTRONIC DRUG LISTING

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AQUA MEDICA, S.A. DE C.V.
Carretera Federal México Cuautla Km. 65.8 No. 8
Tetelcingo, Cuautla 62757
México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**AQUACID-120 Acid Concentrate for Hemodialysis (3,785 mL)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-604-02**
U.S. Agent:	**Latin FDA Registrar**

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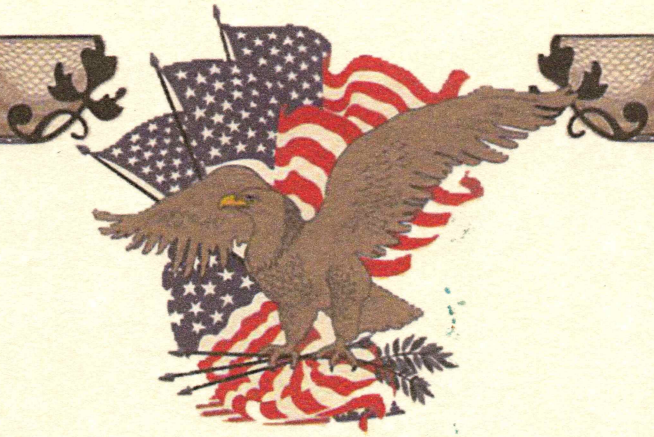

2025-2026

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AQUA MEDICA, S.A. DE C.V.
Carretera Federal México Cuautla Km. 65.8 No. 8
Tetelcingo, Cuautla 62757
México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**AQUACID-200 Acid Concentrate for Hemodialysis (3,785 mL)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-603-03**
U.S. Agent:	**Latin FDA Registrar**

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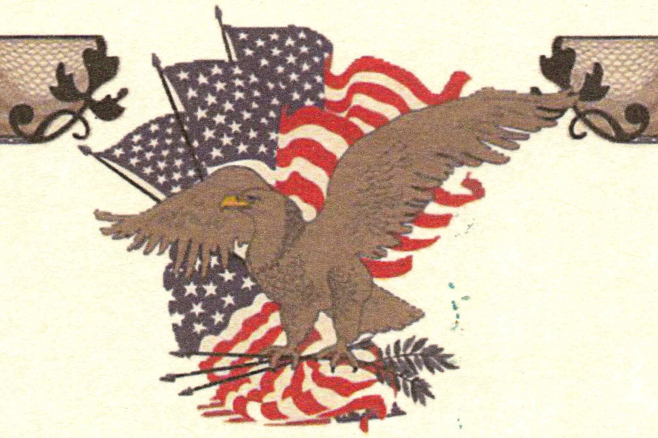

2025-2026

CERTIFICATE OF ELECTRONIC DRUG LISTING

This certifies that:

AQUA MEDICA, S.A. DE C.V.
Carretera Federal México Cuautla Km. 65.8 No. 8
Tetelcingo, Cuautla 62757
México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**AQUACID-220 Acid Concentrate for Hemodialysis (3,785 mL)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-602-04**
U.S. Agent:	**Latin FDA Registrar**

Latin FDA Registrar will confirm that such listing remains effective upon request and presentation of this certificate until the end of the year shown above, unless terminated after issuance of this certificate. Latin FDA Registrar makes no other representations of warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Latin FDA Registrar assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Latin FDA Registrar waives, and will hold harmless and indemnify Latin FDA Registrar from any and all claims in connection with this product, its labeling, FDA drug listing, commerce or use. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of drug listing or the assignment of an DNC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue certificates of drug listing, nor does the U.S. Food and Drug Administration recognize certificates of drug listing. Latin FDA Registrar is not affiliated with the U.S. Food and Drug Administration.

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César López
Executive Director
Latin FDA Registrar Corp.
Dated: November 05, 2025

2025-2026

CERTIFICATE OF REGISTRATION

This certifies that:

AQUA MEDICA, S.A. DE C.V.
Carretera Federal México Cuautla Km. 65.8 No. 8
Tetelcingo, Cuautla 62757
MÉXICO

is registered with the U.S. Food and Drug Administration for the statutory filing period applicable to U.S. FY 2026 pursuant to part 207 of Title 21, U.S. Code of Federal Regulations.

DUNS Number:	**58-969-6442**
Labeler Code:	**81943**
FEI:	**3019430653**
U.S. Agent/Registrant:	**Latin FDA Registrar**

Filling was performed during the October 1 – December 31, 2025 statutory period, and renewal is not required until the next statutory period of October 1 – December 31, 2026. Latin FDA Registrar will confirm that such registration remains effective upon request and presentation of this certificate, until the end of the year stated above, unless terminated after issuance of this certificate. Latin FDA Registrar makes no other representations or warranties, nor does this certificate make any representations of warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Latin FDA Registrar assumes no liability to any person or entity in connection with the foregoing. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of registration or possession of registration number or NDC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Latin FDA Registrar is not affiliated with the U.S. Food and Drug Administration.

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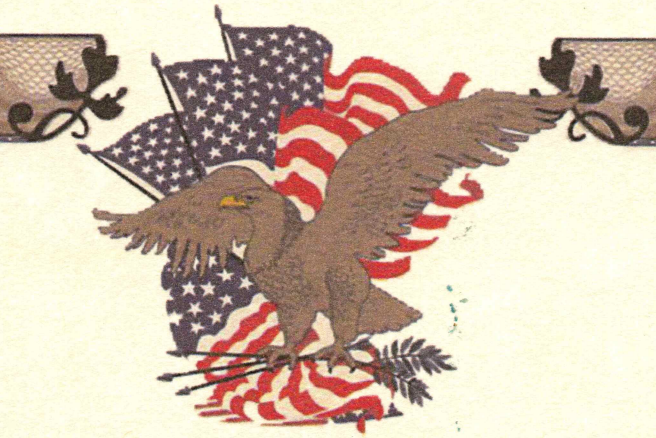

2025-2026

CERTIFICATE OF ELECTRONIC DRUG LISTING

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AQUA MEDICA, S.A. DE C.V.
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Tetelcingo, Cuautla 62757
México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**QUABIC-100 Hemodialysis Grade Sodium Bicarbonate Powder (650 g)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-503-02**
U.S. Agent:	**Latin FDA Registrar**

Latin FDA Registrar will confirm that such listing remains effective upon request and presentation of this certificate until the end of the year shown above, unless terminated after issuance of this certificate. Latin FDA Registrar makes no other representations of warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Latin FDA Registrar assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Latin FDA Registrar waives, and will hold harmless and indemnify Latin FDA Registrar from any and all claims in connection with this product, its labeling, FDA drug listing, commerce or use. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of drug listing or the assignment of an DNC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue certificates of drug listing, nor does the U.S. Food and Drug Administration recognize certificates of drug listing. Latin FDA Registrar is not affiliated with the U.S. Food and Drug Administration.

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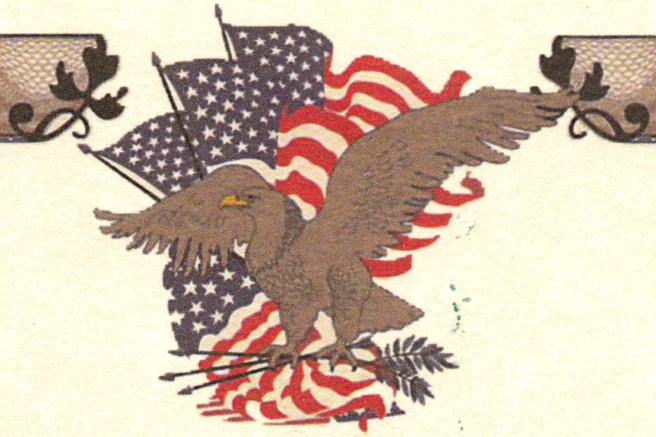

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has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**QUABIC-300 Hemodialysis Grade Sodium Bicarbonate Solution (3,785 mL)**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-502-01**
U.S. Agent:	**Latin FDA Registrar**

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has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**X-TÉRIL GEL Hand Sanitizer 20 L**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-203-03**
U.S. Agent:	**Latin FDA Registrar**

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has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**X-TÉRIL GEL Hand Sanitizer 1000 mL**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-203-01**
U.S. Agent:	**Latin FDA Registrar**

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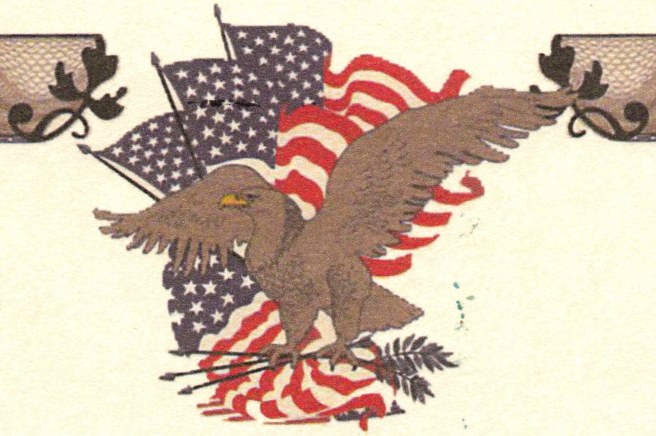

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México

has listed referenced drug product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Latin FDA Registrar:

Product Trade Name:	**X-TÉRIL GEL Hand Sanitizer 3785 mL**
Labeler:	**AQUA MEDICA, S.A. DE C.V.**
Manufacturer:	**AQUA MEDICA, S.A. DE C.V.**
NDC/DLS Numbers:	**81943-203-02**
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